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Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- 1. (Canceled)
- 2. (Previously amended) An isolated polynucleotide comprising nucleotides 3331-3656, 3495-3599 or 3421-3548 of SEQ ID NO: 1 spliced downstream of nucleotides 1-2558 of SEQ ID NO: 1.
 - 3.-21. (Canceled)
- 22. (Previously amended) An isolated polynucleotide comprising a smooth muscle (SM) α-A promoter/enhancer in operable association with a heterologous polynucleotide, wherein the promoter/enhancer comprises sufficient sequence from the first intron of the SM α-A gene to confer smooth muscle cell-specific expression *in vivo* and wherein the promoter/enhancer hybridizes to the complement of SEQ ID NO:1 when DNA comprising the complement of SEQ ID NO:1 is hybridized in 0.5 M NaHPO4, 7% sodium dodecyl sulfate (SDS), 1mM EDTA at 65° C, and washed in 0.1xSSC/0.1% SDS at 68° C.

23-30. (Canceled.)

31. (Currently amended) The isolated polynucleotide of claim 22, wherein the sequence from the first intron comprises the rat AP1-like, Int CArG and GATA elements, wherein

the AP1-like element comprises SEQ ID NO:19 with the following change: T22C SEQ ID NO:26;

the Int CArG element comprises SEQ ID NO:16; and

the GATA element comprises SEQ ID NO:20 with the following changes: G12T, G14A and C18T SEQ ID NO:31.

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- 32. (Previously amended) The isolated polynucleotide of claim 22, wherein the sequence from the first intron comprises SEQ ID NO:8.
- 33. (Previously amended) The isolated polynucleotide of claim 22, wherein the promoter/enhancer comprises the CArG B and CArG A elements depicted in SEQ ID NO:15 and SEQ ID NO:14, respectively.
- 34. (Previously amended) The isolated polynucleotide of claim 22, wherein the promoter/enhancer comprises the sequence depicted in SEQ ID NO:4.
- 35. (Previously presented) A vector comprising the polynucleotide of claim 22.
- 36. (Previously amended) An isolated genetically-engineered host cell comprising a polynucleotide comprising a SM α-A promoter/enhancer in operable association with a heterologous polynucleotide, wherein the promoter/enhancer comprises sufficient sequence from the first intron of the SM α-A gene to confer smooth muscle cell-specific expression *in vivo* and wherein the promoter/enhancer hybridizes to the complement of SEQ ID NO:1 when DNA comprising the complement of SEQ ID NO:1 is hybridized in 0.5 M NaHPO4, 7% sodium dodecyl sulfate (SDS), 1mM EDTA at 65° C, and washed in 0.1xSSC/0.1% SDS at 68° C.

37-40. (Canceled)

41. (Previously amended) The host cell of claim 36, wherein the sequence from the first intron comprises the rat AP1-like, Int CArG and GATA elements wherein the AP1-like element comprises SEQ ID NO:19 with the following change: T22C SEQ ID NO:26;

the Int CArG element comprises SEQ ID NO:16; and the GATA element comprises SEQ ID NO:20 with the following changes: G12T, G14A and C18T SEQ ID NO:31.

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- 42. (Canceled)
- 43. (Previously presented) The host cell of claim 36, wherein the promoter/enhancer comprises the nucleotide sequence of SEQ ID NO:1.
- 44. (Previously presented) The host cell of claim 36, wherein the sequence from the first intron comprises SEQ ID NO:8.
- 45. (Previously presented) The host cell of claim 36, wherein the promoter/enhancer comprises the CArG B and CArG A elements depicted in SEQ ID NO:15 and SEQ ID NO:14, respectively.
- 46. (Previously presented) The host cell of claim 36, wherein the promoter/enhancer comprises the sequence depicted in SEQ ID NO:4.
- 47. (Previously presented) The isolated polynucleotide of claim 22, wherein the promoter/enhancer comprises nucleotides 1-2605, 2011-2605, 2011-5342, 3331-3656, 3421-3548 or 3495-3599 of SEQ ID NO:1.
- 48. (Previously presented) The isolated polynucleotide of claim 22, wherein the promoter/enhancer comprises the nucleotide sequence of SEQ ID NO:1.